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# Faceting and stability of smectic A droplets on a solid substrate

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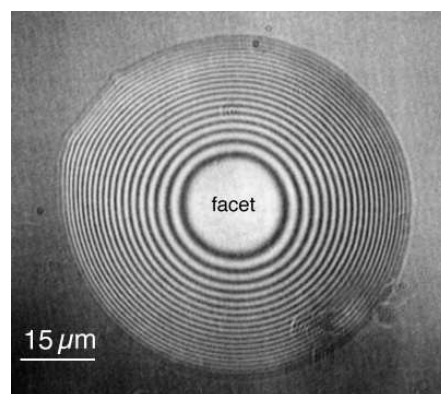
**Abstract.** It is shown that a smectic A droplet deposited on a solid substrate treated for strong homeotropic anchoring is faceted at the top in spite of the fact that there are no steps at the free surface, but instead edge dislocations in the bulk. The radius of the facet and the full profile of the curved part of the droplet are determined as a function of the temperature in the vicinity of a nematic-smectic A phase transition. It is shown that the observed profiles do not correspond to the actual equilibrium shape, but to metastable configurations close to their point of marginal stability. In addition, we predict that the profiles must be different for a given temperature depending on whether the droplet has been heated or cooled down to reach this temperature. Finally, we discuss the problem of the formation of giant dislocations in big droplets (Grandjean terraces).

**PACS.** 68.03.Cd Surface tension and related phenomena – 61.30.Hn Surface phenomena: alignment, anchoring, anchoring transitions, surface-induced layering, surface-induced ordering, wetting, prewetting transitions, and wetting transitions – 61.30.Jf Defects in liquid crystals

## 1 Introduction

The equilibrium shape of a crystal is obtained by minimizing its total surface energy. This procedure leads to the well-known Wulff construction according to which the equilibrium shape is the pedal of the polar plot of the surface energy  $\gamma(\theta, \varphi)$  (with  $(\theta, \varphi)$  the angles defining the orientation of the surface element with respect to the underlying crystal lattice [1,2]). In three-dimensional crystals, this construction leads to faceted shapes, with the size and the number of facets decreasing when the temperature increases. Such shapes were observed by scanning electron microscopy in micron-sized crystals of ordinary metals such as lead [3], silicon [4], or silver [5]. Large He<sup>4</sup> crystals (*i.e.*, of centimetric size) are also faceted, with the advantage of equilibrating much faster than the former due to their quantum nature (atoms can go through energy barriers by tunneling effect) [6]. Faceted shapes were also observed in soft matter, in particular in the Blue Phases of cholesteric liquid crystals [7–9], in the smectic B plastic crystals [10,11], and in the cubic phases of the lyotropic systems [12–15]. Although we are dealing with monocrystals in all these examples, the real question is always to know whether the shapes observed experimentally are at equilibrium (in a stable or a metastable state), or result from some growth processes. This delicate problem will be discussed in detail in this article.

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**Fig. 1.** Observation in Michelson interferometry of a faceted droplet of the liquid crystal 8CB placed on a glass slide treated for homeotropic anchoring (photograph by J. Bechhoefer, 1991).

Faceted droplets were also observed in smectic A liquid crystals which are one-dimensional crystals (the rod-like molecules form fluid layers which pile up on each other). In that case, the droplets are deposited on solid substrates treated for homeotropic anchoring. Although first observations of faceting in smectic A are old [16–18], precise determination of droplet profiles as a function of temperature close to a nematic phase or an isotropic liquid is

relatively recent [19,20]. The main findings of these experimental studies can be summarized as follows:

1. Each droplet shows a single facet parallel to the substrate (Fig. 1), whose size increases when the temperature decreases.

2. The curved part of the droplet surface joins tangentially onto the facet.

3. All the attempts to fit the experimental profiles to a power law of type  $(\rho - r_f)^\nu$ , with  $\rho$  the radius in polar coordinates and  $r_f$  the facet radius, led to an exponent  $\nu$  ranging typically between 1 and 2.5, depending on the temperature and the “angular window” of the fit. These large variations suggest that a power law is certainly not a good choice.

4. For a given temperature, the facet radius depends on the thermal history of the droplet. More precisely, the facet is systematically larger after heating than after cooling from the high-temperature phase (*i.e.*, the nematic phase or the isotropic liquid, depending on the material).

5. Finally, the facet size stabilizes much faster after cooling (within a few seconds or minutes, typically) than after heating: in that case, many hours or even days are necessary to reach a stable shape with a facet generally larger than that observed by direct cooling from the nematic or the isotropic liquid.

To explain these observations, we used in ref. [21] the terrace-ledge-kink model (or TLK model) according to which the layers are strictly parallel to the solid surface and emerge at the free surface by forming steps. This classical model predicts that a facet forms at the top of the droplet and matches tangentially the curved part of the droplet surface. This particular matching comes from the fact that the steps repulse with a  $1/d^2$ -law, which, in smectic A, is only due to their entropic interactions ( $d$  is the distance between two steps), their elastic interactions being completely negligible because of the fluidity of the layers. This generic interaction leads to profiles of type  $(\rho - r_f)^\nu$ , with the universal exponent  $\nu = 3/2$ , in profound disagreement with experiments.

In the same article, we developed an alternative model, in which steps were replaced by bulk dislocations. A direct consequence was that the top smectic layer was bent parallel to the free surface. It turns out that this model was more reasonable than the TLK model because we know both theoretically [22] and experimentally (for a review, see [11]) that in smectic A, edge dislocations are repulsed by the free surface, a point we detail in reference [23]. On the other hand, we assumed without any justification, that the droplet had the shape of a spherical cap. In this article, we show that this affirmation was wrong and that this model can explain as well the existence of faceted droplets.

The plan of the article is as follows. In Section 2, we recall the basics of the model and calculate the droplet energy and its general profile. In Section 3, we show the existence of two facet radii corresponding to two extrema of the total energy. In Section 4, we demonstrate that the smaller radius corresponds to the critical radius of the facet while the other, twice larger, corresponds to the equilibrium radius of the facet. We also show that all

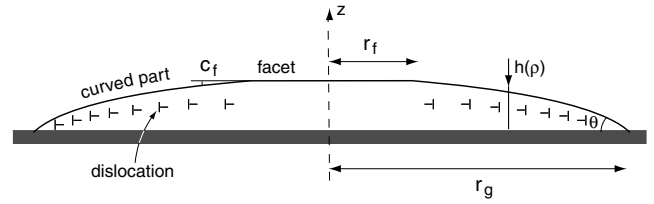


Fig. 2. Droplet configuration.

the droplets whose facet radius is larger than the critical radius are metastable. We thus conclude that the facet radius measured experimentally after cooling down the droplet from the nematic or the isotropic phases is the critical radius. In Section 5, we calculate the profile of a droplet at the critical radius as a function of the temperature close to a second-order smectic A-nematic phase transition and we compare with experiments. We then analyze in Section 6 how the facet radius changes in time when the temperature is again increased. A criterion of marginal stability is proposed to explain why the facet radius again increases in spite of the fact that the droplet height is blocked. In Section 7, we discuss the stability of the droplets with respect to the formation of giant dislocations. This mechanism allows us to explain the formation of the so-called Grandjean terraces in big droplets. Finally, some concluding remarks are drawn in Section 8.

## 2 Energy of a faceted droplet and general surface profile

The droplet configuration is recalled in Figure 2. It has the circular symmetry around the  $z$ -axis perpendicular to the substrate treated for homeotropic anchoring. Because the dislocations are repelled by both the substrate and the free surface, the dislocations are distributed throughout the bulk in order to accommodate the macroscopic faceted shape. In the following, we denote by  $r_g$  the droplet radius at the substrate, by  $r_f$  the facet radius, by  $c_f$  the matching angle between the facet and the curved part of the droplet surface, and by  $V$  the droplet volume, supposed to be constant.

In order to calculate the macroscopic droplet profile  $h(\rho)$  (with  $\rho$  the radius in polar coordinates), we need to minimize the total energy  $e$  of the droplet. The latter contains two terms:

1. A surface term coming from the different surface energies  $\gamma$ ,  $\gamma_{SL}$ , and  $\gamma_{SA}$  between, respectively, the liquid crystal and the air, the substrate and the liquid crystal, and the substrate and the air.
2. A bulk term due to the presence of the dislocations, which roughly sit at mid-distance between the solid and the free surfaces, as shown in reference [23].

Let  $E_d$  be the energy of a dislocation per unit length and  $b$  its Burgers vector (which we assume to be equal to the layer thickness  $d$  for the moment). A straightforward calculation, similar to that already used to calculate the profile of a meniscus surrounding a free-standing

film [11, 24], yields:

$$e = 2\pi \int_0^{r_g} \rho d\rho (\gamma_{SL} - \gamma_{SA}) + 2\pi\gamma \int_0^{r_f} \rho d\rho + 2\pi\gamma \int_{r_f}^{r_g} \rho d\rho \sqrt{1 + \left(\frac{dh}{d\rho}\right)^2} - 2\pi \frac{E_d}{b} \int_{r_f}^{r_g} \rho \left(\frac{dh}{d\rho}\right) d\rho. \quad (1)$$

In this formula, the first term gives the surface energy on the substrate, the next two the surface energy between the droplet and the air, and the fourth one the energy of the bulk dislocations (with  $-(1/b)(dh/d\rho)$  representing the density of dislocations). It is important to note that in this formula, the free surface of the droplet is assumed to be smooth at the scale of the distance between dislocations. This is justified in a droplet because the dislocations sit at large distance from the free surface (see reference [23]), so that their deformation fields strongly overlap close to the free surface. For that reason, the fourth term in equation (1) in  $\sqrt{1 + (dh/d\rho)^2}$  may be considered as the repulsive interaction term between the dislocations.

In practice, the droplets are rather flat (the wetting angle at the solid substrate is usually less than  $15^\circ$ ). For that reason, we can reasonably assume that  $|dh/d\rho| \ll 1$ , which allows us to develop the term  $\sqrt{1 + (dh/d\rho)^2}$  to first order in  $(dh/d\rho)^2$ . This leads to the following, simpler, expression for the energy:

$$e \approx 2\pi S \int_0^{r_g} \rho d\rho + \pi\gamma \int_{r_f}^{r_g} \rho d\rho \left(\frac{dh}{d\rho}\right)^2 - 2\pi \frac{E_d}{b} \int_{r_f}^{r_g} \rho \left(\frac{dh}{d\rho}\right) d\rho, \quad (2)$$

where  $S = \gamma_{SL} - \gamma_{SA} + \gamma$  is the so-called spreading power.

The following step consists of minimizing the energy with the constraint that the volume remains constant:

$$V = 2\pi \int_0^{r_g} \rho h(\rho) d\rho \quad (3)$$

This leads to the following differential equation for the profile (equivalent to the Laplace law):

$$\Delta P = -\gamma \frac{1}{\rho} \frac{d}{d\rho} \left[ \rho \frac{dh}{d\rho} \right] + \frac{E_d}{b\rho}, \quad (4)$$

where  $\Delta P = P_L - P_A$  is the pressure difference between the liquid crystal and the air. Note that  $\Delta P$  is the Lagrange multiplier associated with condition (3).

The resolution of equation (4) leads to the following general profile for the droplet:

$$h(\rho) = \frac{E_d}{b\gamma} \left( \rho - r_g - r_f \ln \frac{\rho}{r_g} \right) - \frac{\Delta P}{2\gamma} \left( \frac{\rho^2 - r_g^2}{2} - r_f^2 \ln \frac{\rho}{r_g} \right) - c_f r_f \ln \frac{\rho}{r_g}. \quad (5)$$

This profile satisfies the conditions  $h(r_g) = 0$  and  $[dh(\rho)/d\rho]_{\rho=r_f} = -c_f$ , with  $c_f > 0$ .

Although we know now the general droplet profile, the problem is far from being solved because  $r_g$ ,  $r_f$ , and  $c_f$  are still unknown.

Determining these quantities requires to minimize the total energy. The latter can be calculated by replacing  $h(\rho)$  by its expression (5) in equation (2). This leads to the following formula for the energy:

$$e = e_o + \alpha c_f + \beta c_f^2 \quad (6)$$

with

$$e_o = \frac{\pi\gamma r_g^2}{2} \left[ s - g(g - pr_f) \frac{(r_g^2 - r_f^2)}{r_g^2} + \frac{(pr_f)^2}{8} \left( 3 \frac{r_f^2}{r_g^2} - 4 + \frac{r_g^2}{r_f^2} \right) + \frac{r_f^2}{2r_g^2} (2g - pr_f)^2 \ln \frac{r_g}{r_f} \right], \quad (7a)$$

$$\alpha = \frac{\pi\gamma r_f}{2} \left[ p(r_g^2 - r_f^2) + 2r_f(2g - pr_f) \ln \frac{r_g}{r_f} \right], \quad (7b)$$

$$\beta = (\pi\gamma r_f^2) \ln \frac{r_g}{r_f}. \quad (7c)$$

In these equations we have introduced the following notations:

$$g = \frac{E_d}{b\gamma}, \quad s = \frac{2S}{\gamma}, \quad \text{and} \quad p = \frac{\Delta P}{\gamma}. \quad (8)$$

Note that  $g$  and  $s$  are dimensionless constants characteristic of the material and that  $p$  (which gives the pressure difference between the interior of the droplet and the air) has for dimension  $[L]^{-1}$ .

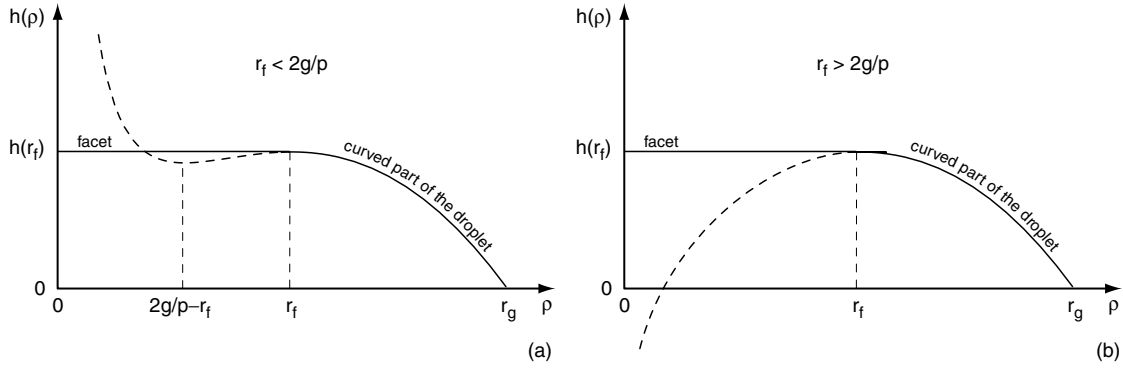
### 3 Determination of the facet and the droplet radii

Before minimizing the energy  $e$  as a function of  $r_g$  and  $r_f$ , let us show that  $c_f = 0$ .

According to equation (7c),  $\beta > 0$  because  $r_g > r_f$ . The sign of  $\alpha$  is less obvious to determine. Nevertheless it will be possible to check *a posteriori* that  $p > 0$  (the pressure inside the droplet is always larger than the air pressure). In this case,  $\alpha > 0$  according to equation (7b) whatever the value of  $r_f < r_g$ . As a result, the energy  $e$  is minimum when  $c_f = 0$  and reduces to the simplified expression  $e_o$  given by equation (6).

Another remark concerns the droplet profile  $h(\rho)$  given by equation (5) when we impose  $c_f = 0$ . Indeed, this function has two extrema given by  $\rho_1 = r_f$  and  $\rho_2 = 2g/p - r_f$  (Fig. 3). It can be easily checked that the profile is physically acceptable if  $\rho_2 < \rho_1$ , *i.e.* if  $r_f \geq g/p$ . We will show in Section 4 that this condition is indeed fulfilled experimentally.

Let us now minimize  $e_o$  as a function of the two radii  $r_f$  and  $r_g$ . Note that in the expression (7a) of  $e_o$ ,  $p$  is also a function of  $r_f$  and  $r_g$  given by the condition of fixed



**Fig. 3.** Typical plots of the function  $h(\rho)$  when  $g/p < r_f < 2g/p$  (a) and  $r_f > 2g/p$  (b). In both cases, the dashed part of the curve has no physical meaning.

volume:

$$V = \pi r_f^2 h(r_f) + 2\pi \int_{r_f}^{r_g} \rho h(\rho) d\rho = \frac{\pi r_g^3}{24} (1-k)^2 (3pr_g(1+k)^2 - 4g(2+k)) \quad (9)$$

with  $k = r_f/r_g$ . As a consequence, minimization with respect to  $r_f$  reads:

$$\frac{\partial e_o}{\partial r_f} + \frac{\partial e_o}{\partial p} \frac{\partial p}{\partial r_f} = 0. \quad (10)$$

This equation has two simple solutions:

$$r_f = \frac{g}{p} \quad \text{or} \quad r_f = \frac{E_d}{b\Delta P} \quad (11a)$$

and

$$r_f = \frac{2g}{p} \quad \text{or} \quad r_f = \frac{2E_d}{b\Delta P}, \quad (11b)$$

where the pressure  $p$  is given by equation (9) as a function of  $V$  and  $r_g$ .

Minimization of  $e_o$  with respect to  $r_g$  reads similarly

$$\frac{\partial e_o}{\partial r_g} + \frac{\partial e_o}{\partial p} \frac{\partial p}{\partial r_g} = 0. \quad (12)$$

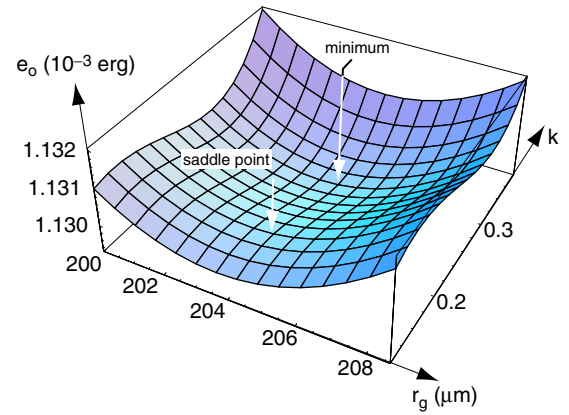
This equation may be expressed as a function of  $k$  using Mathematica. For the first solution (11a), it reads explicitly

$$(1+k) \left( -1+3k-4k^2+k^4+k^5+4s \left( \frac{k}{g} \right)^2 (1+k) \right) = 8k^4 \ln k, \quad (13a)$$

while for solution (11b),  $k$  has a simple expression,

$$k = \frac{g}{g + \sqrt{s}}. \quad (13b)$$

In conclusion, the energy  $e_o$  has two extrema which are given (for a droplet of a given volume  $V$ ) by equations (9,



**Fig. 4.** 3D plot of the droplet energy  $e_o$  as a function of  $k = r_f/r_g$  and  $r_g$ . The saddle point and the absolute minimum are indicated by two arrows (8CB,  $T = 24^\circ\text{C}$ ,  $V = 10^{-6} \text{ cm}^3$ ).

11a), and (13a) and by equations (9, 11b), and (13b), respectively.

The problem that now arises is to determine the nature of these two solutions. To answer this question, we plotted the energy  $e_o$  as a function of  $r_f$  and  $r_g$ , or equivalently as a function of  $k$  and  $r_g$ . An example is shown in Figure 4 for a droplet of volume  $V = 10^{-6} \text{ cm}^3$  and typical values of the material parameters (corresponding to the liquid crystal 8CB at  $T = 24^\circ\text{C}$ ):  $\gamma = 31 \text{ erg/cm}^2$  [25],  $g = 0.073$  [26] and  $s = 0.0304$  (this value corresponds to a wetting angle  $\theta$  of  $10^\circ$  in the nematic phase, knowing that in that case  $s = 2(1 - \cos \theta)$ ). This plot shows that the energy possesses a saddle point corresponding to the first solution (11a) and a true minimum corresponding to the second solution (11b). As a consequence, solution (11b) gives the equilibrium configuration, whereas solution (11a) is unstable. This graph also shows that there is a valley joining these two solutions. More precisely, for each value of  $k$  lying between  $g/(pr_g)$  and  $2g/(pr_g)$ , the energy passes through a minimum as a function of  $r_g$ . The question we tackle in the next section is to determine whether these intermediate solutions are accessible experimentally and observable over long time (which would suggest they are metastable).

#### 4 On the metastability of faceted droplets

To answer this question, the global method developed in the previous sections is not adequate. A more local approach is necessary. In this respect, let us consider the local mechanical equilibrium of the layers at the free surface. At radius  $\rho$ , this equation reads:

$$\Delta P - \sigma = \gamma C, \quad (14a)$$

where  $C$  is the local curvature of the surface and  $\sigma$  the elastic stress normal to the layers. Expressing  $C$  as a function of  $h(\rho)$  gives in the curved part of the droplet:

$$\Delta P = -\gamma \frac{1}{\rho} \frac{d}{d\rho} \left[ \rho \frac{dh}{d\rho} \right] + \sigma. \quad (14b)$$

By comparing equation (4) with equation (14b), we obtain the following equation, valid in the curved part of the droplet:

$$\sigma(\rho > r_f) = \frac{E_d}{b\rho}. \quad (15)$$

This equation expresses the mechanical equilibrium of the dislocations inside the droplet. Indeed,  $\sigma(\rho)b$  is the Peach and Koehler force acting on the dislocation of radius  $\rho$ , while  $E_d/\rho$  is the line tension force. These two forces equilibrate, the former tending to grow the dislocation whereas the latter tends to shrink it.

Under the facet, which is flat, equation (14) becomes

$$\sigma(\rho < r_f) = \Delta P. \quad (16)$$

As a consequence, the stress is generally different from each side of the facet edge. Note that a similar situation exists in a free-standing smectic film surrounded by a straight meniscus. In that case,  $\sigma = \Delta P$  in the film, whereas  $\sigma = 0$  in the meniscus [11,24].

Let us now compare the stress from each side of the facet edge. According to equations (15) and (16), we have:

$$\sigma(\rho = r_f^+) = \sigma^+ = \frac{E_d}{br_f} \quad (17a)$$

whereas

$$\sigma(\rho = r_f^-) = \sigma^- = \Delta P. \quad (17b)$$

These two equations show that  $\sigma^+ = \sigma^-$  at  $r_f = E_d/(b\Delta P)$ . This radius corresponds to the saddle point of the total energy  $e_o$ , *i.e.* to the solution given by equations (11a) and (13a). In the following, we will denote by  $r_c$  this particular solution. We immediately see from equation (17) that if  $r_f < r_c$ , then  $\sigma^- < \sigma^+$ . As a consequence the dislocation which borders the facet collapses and the facet grows. For that reason,  $r_c$  corresponds to the *critical radius* of the facet.

Let now consider a droplet with a facet of radius  $r_f > r_c$ . In this particular case,  $\sigma^- > \sigma^+$ . As a consequence the dislocation bordering the facet will not collapse, except if

the thermal fluctuations (*i.e.*, the external medium) brings enough energy to overcome the energy barrier

$$W^- = \int_{r_f}^{r_c} \left( \frac{E_d}{\rho} - \Delta P b \right) 2\pi \rho d\rho = 2\pi b \Delta P \int_{r_f}^{r_c} (r_c - \rho) d\rho = \pi b \Delta P (r_f - r_c)^2. \quad (18)$$

In practice,  $W^- \gg k_B T$  as long as the facet radius is a bit larger than the critical radius. To fix ideas, let us consider the example of Figure 4 which would correspond to a droplet of 8CB at 24 °C of volume  $V = 10^{-6} \text{ cm}^3$  and external radius  $r_g \approx 204 \mu\text{m}$ : in that case, we calculate  $r_c = 32 \mu\text{m}$ ,  $\Delta P \approx 730 \text{ dyn/cm}^2$ , which gives  $W^- \approx 7 \times 10^{-13} \text{ erg} \approx 80 k_B T$  for a facet radius  $r_f = 1.02 r_c$ .

This simple calculation shows that, in practice, the facet radius which is measured experimentally after cooling down the droplet from the high-temperature phase must be very close to the critical radius.

One can also calculate the typical time a droplet needs to reach its new configuration when the temperature is decreased. Indeed, let us consider a droplet with a facet radius larger than the critical radius. The dislocation close to the facet is unstable and tends to collapse. Indeed, it experiences three forces: the inward force  $-E_d/\rho$  associated with the line tension  $E_d$  and two outward forces, the Peach and Koehler force of amplitude  $\Delta P b$  and a frictional one, proportional to the dislocation velocity and of expression  $-(b/m)(d\rho/dt)$ , where  $m$  is dislocation mobility [11]. Equating these three forces, we have

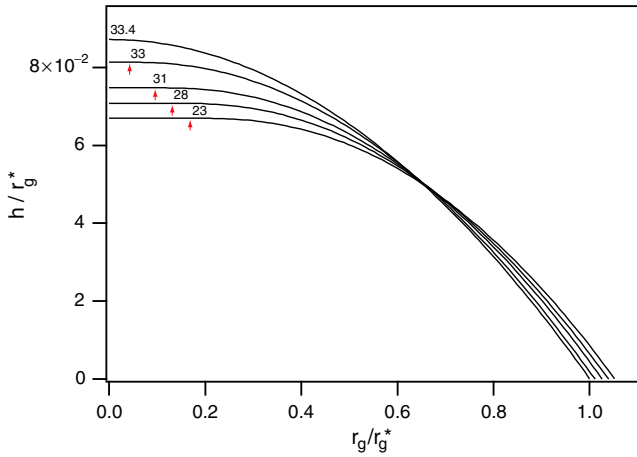
$$\frac{b}{m} \frac{d\rho}{dt} = \Delta P b - \frac{E_d}{\rho} = \Delta P b \left( 1 - \frac{r_c}{\rho} \right). \quad (19)$$

As expected, this equation states that the innermost dislocation collapses as  $r_f$  is smaller than  $r_c$ . Its collapse time can be calculated from this equation (for the calculation, see [11], p. 450) and reads:

$$\tau_{\text{coll}} = \frac{r_c}{m\Delta P} \left[ \ln \left( \frac{1}{1-x} \right) - x \right] = \tau \left[ \ln \left( \frac{1}{1-x} \right) - x \right]. \quad (20)$$

In this expression,  $x = r_f/r_c < 1$  and  $\tau = r_c/(m\Delta P)$  fixes the time scale. In 8CB,  $m$  is of the order of  $3 \times 10^{-7} \text{ cm}^2 \text{ sg}^{-1}$  at 24 °C [11]. For a droplet of typical volume  $V = 10^{-6} \text{ cm}^3$ , we calculated in the previous paragraph  $r_c \approx 32 \mu\text{m}$  and  $\Delta P \approx 730 \text{ dyn/cm}^2$ , which yields  $\tau = 15 \text{ s}$ . This time fixes the typical equilibration time of the droplet (for  $x = 0.98$ ,  $\tau_{\text{coll}} \approx 3\tau$ ). We emphasize that the latter is certainly much shorter in the experiments cited in references [19,20] because the droplets studied had smaller volumes (so that  $r_c$  was smaller and  $\Delta P$  larger) and the temperature was higher (4O8 and 8OCB melt at higher temperatures than 8CB). We thus expect the dislocation mobility  $m$  be larger than  $3 \times 10^{-7} \text{ cm}^2 \text{ sg}^{-1}$  in these materials, inasmuch as this quantity is thermally activated and strongly increases when the temperature increases [11].

This calculation thus explains why the droplets rapidly reach their critical radii when the temperature is decreased.



**Fig. 5.** Four normalized profiles calculated at four different temperatures for the liquid crystal 8CB. The temperature is given beside each profile. An arrow marks the edge of the facet on each profile. At 33.4 °C, the droplet has the shape of a spherical cap.

In the following section, we show the temperature evolution of the droplet profile when  $r_f = r_c$  in the vicinity of a second-order nematic-smectic A phase transition.

## 5 Evolution of the droplet profile and of the facet radius at decreasing temperature (i.e., when $r_f = r_c$ )

In order to find the droplet profile at the critical radius, we need to solve the set of equations (9, 11a), and (13a). More precisely, equation (13a) gives  $k$ . Then, we calculate  $pr_g = g/k$  from equation (11a), which gives after substitution into equation (9):

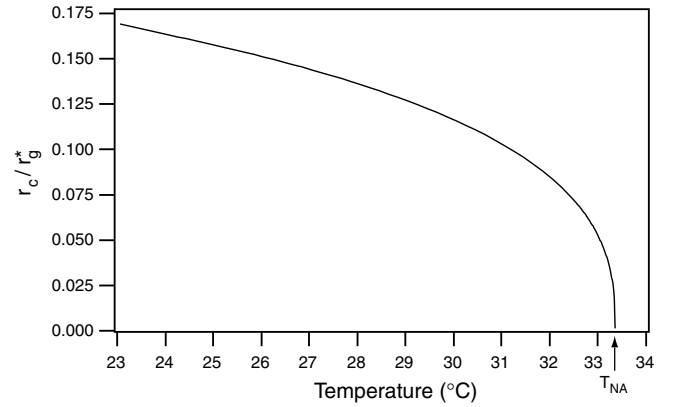
$$V = \frac{\pi r_g^3}{24} g(1-k)^2(3(1+k)^2/k - 4(2+k)). \quad (21)$$

This equation allows us to calculate  $r_g$  for a given volume  $V$ . As for the droplet profile, it is given by equation (5) by taking  $c_f = 0$ .

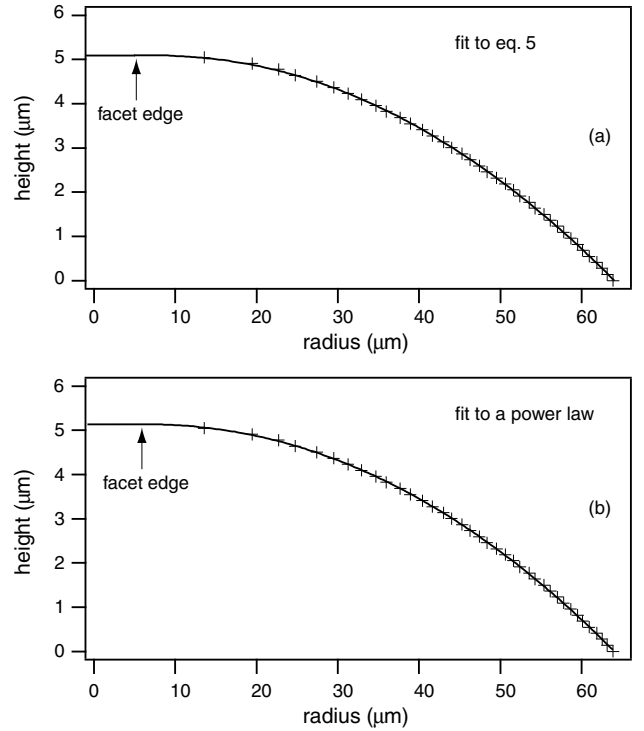
In practice, the shapes are self-similar, so it is preferable to use as unit length the radius  $r_g^*$  of the droplet in the nematic phase. The droplet is then at equilibrium and has the shape of a spherical cap with a contact angle at the substrate given by the Young equation:  $s = 2(1 - \cos \theta)$ . As for its volume, it may be simply obtained by making  $k \rightarrow 0$  in equations (13a) and (21). We successively obtain  $g/k = 2\sqrt{s}$  and  $V = (\pi r_g^{*3}/8)(g/k)$ , from which we calculate

$$r_g^* = \left( \frac{4V}{\pi\sqrt{s}} \right)^{1/3}. \quad (22)$$

Figure 5 shows five profiles (normalized to  $r_g^*$ ) calculated for the liquid crystal 8CB at four different temperatures:  $T = T_{NA} = 33.4$  °C (transition temperature, spherical cap),  $T = 33$  °C,  $T = 31$  °C,  $T = 28$  °C, and  $T = 23$  °C. As expected, the droplet height decreases due



**Fig. 6.** Normalized critical radius as a function of the temperature calculated for the liquid crystal 8CB.



**Fig. 7.** a) Experimental profile (crosses) determined from the photograph shown in Figure 1. The solid line is the best fit to equation (5) with  $c_f = 0$ . The arrow marks the facet edge given by the fit. The latter is pretty far from the last fringe detected experimentally. This shows all the difficulty to measure the facet radius (the top of the droplet is very flat). b) Same experimental data fitted to a power law. The size of the facet is found a bit larger than using the previous fit and the profile exponent is  $\nu = 2.1$ .

to the rapid collapse of the inner dislocations and  $r_g$  increases when the temperature decreases.

Figure 6 shows the normalized critical facet radius  $r_c$  as a function of temperature. The latter vanishes at the transition temperature, in agreement with experiments [19,20]. Note that all the calculations were

performed using  $E_d/b = 0.84(33.4 - T)^{0.446}$  (in CGS unit) [26] and by assuming that  $b$ ,  $\gamma$ , and  $s$  are independent of the temperature ( $b = 30 \text{ \AA}$ ,  $\gamma = 31 \text{ erg/cm}^2$  [25], and  $s = 0.0304$ , which corresponds to a contact angle  $\theta = 10^\circ$  when the droplet has a spherical shape).

To conclude this section, let us compare these predictions with experiments. Figure 1 shows a droplet of 8CB deposited on a glass plate treated with a silane for homeotropic anchoring. The photograph was taken at  $30^\circ\text{C}$  by John Bechhoefer in 1991 using a Michelson interferometer. In this picture, the fringes give a contour map with  $\lambda/2 \approx 273 \text{ nm}$  separating each fringe. An experimental profile obtained by detecting the positions of the fringes along a diameter is shown in Figure 7a (crosses).

This profile was fitted to equation (5) (with  $c_f = 0$ ) by using  $E_d/(b\gamma)$ ,  $\Delta P/(2\gamma)$ , and the radius of the facet  $r_f$  as fit parameters (with the droplet radius  $r_g$  being known). We found that the fit is excellent over the whole profile. The values of the fit parameters obtained from Igor Pro are  $E_d/(b\gamma) = 0.035$ ,  $\Delta P/(2\gamma) = 34 \text{ cm}^{-1}$ , and  $r_f = 5 \mu\text{m}$  (for a droplet radius  $r_g = 64 \mu\text{m}$ ), which gives  $E_d \approx 3.2 \times 10^{-7} \text{ dyn}$  for 8CB at  $30^\circ\text{C}$ . This value is in reasonable agreement with that calculated from formula  $E_d/b = 0.84(33.4 - T)^{0.446}$  given in reference [26] (which gives  $E_d \approx 4.3 \times 10^{-7} \text{ dyn}$ ). This point is important to justify the model. In addition, it must be emphasized that the fit of the curved part of the droplet gives also the value of the critical radius. Indeed, according to equation (11a),  $r_c = E_d/(b\Delta P)$ , which yields by using the fit parameters:  $r_c = 0.035/(2 \times 34) \text{ cm} \approx 5.1 \mu\text{m}$ . We thus conclude that for the droplet shown in Figure 1,  $r_f = r_c$  within experimental errors. This is direct evidence that we measure the critical radius instead of the true equilibrium radius, when measurements are performed after cooling down the droplet.

For comparison, we fitted the same data with a power law of type  $h(\rho) = h_f - \varpi(\rho - r_f)^\nu$ . As shown in Figure 7b, the fit is as good as with equation (5) and gives  $h_f = 5.1 \mu\text{m}$ ,  $\varpi = 25$  (in CGS units),  $r_f = 5.8 \mu\text{m}$  and  $\nu = 2.09$ . We note that the facet size obtained this way is comparable with that found by using equation (5) ( $5.8 \mu\text{m}$  instead of  $5.1 \mu\text{m}$ ). As for the exponent  $\nu$ , it is found close to 2 (the same value was measured in 8OCB and 4O8 close to the nematic-smectic A transition temperature [19,20]). Within the framework of the TLK model, such a value of  $\nu$  suggests that the droplet is not at equilibrium, but at its critical radius [2,19]. From this point of view, the two models lead to the same conclusion.

We nevertheless prefer the present model because it is now well established from experiments on free-standing films (for a review, see Chapt. 8 in Ref. [11]) that dislocations are strongly repulsed from the free surface in smectic A liquid crystals. In addition, this model applies along the whole profile by construction, contrary to the TLK model, which is only valid in the vicinal regime (*i.e.*, close to the facet where steps are well separated).

## 6 Facet radius and profile evolution at increasing temperature

Let us now consider a droplet at low temperature obtained after decreasing the temperature from the nematic phase. We have shown that its facet radius is given by equation (11a), which corresponds to the critical radius  $r_c$ . In the following we denote by  $h_f$  the height of the droplet at this temperature.

The delicate problem we would like to tackle in this section is the following: what happens if the temperature is again increased. The droplet is then out of equilibrium since its profile no longer satisfies the equilibrium equation (4). So its shape must slowly change (slowly, because any deformation of the profile implies a redistribution of the dislocations, which is strongly dissipative).

To fix ideas, let us start from the profile shown in Figure 5 at  $T_i = 23^\circ\text{C}$  and suppose we increase the temperature of the droplet up to  $T_f = 33^\circ\text{C}$ . Will the droplet recover the shape shown in Figure 5 at this temperature?

To reach again the final state described in Figure 5, dislocation loops must nucleate since the droplet height must increase. But the nucleation of a dislocation loop imposes to overcome an energy barrier of height

$$W^+ = \int_0^{r_c} \left( \frac{E_d}{\rho} - \Delta P b \right) 2\pi\rho d\rho = 2\pi b \Delta P \int_0^{r_c} (r_c - \rho) d\rho = \pi b \Delta P r_c^2. \quad (23)$$

A rapid numerical estimate shows that  $W^+$  is always much larger than  $k_B T$  in experiments, so nucleation is forbidden. The direct consequence is that the droplet cannot recover its initial shape.

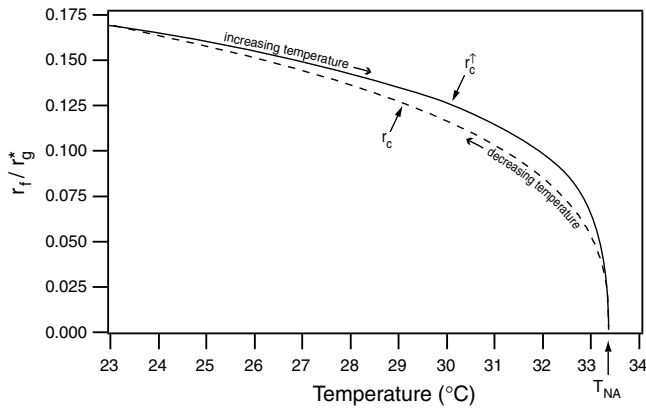
This conclusion (the same was drawn in Ref. [19] from the TLK model) is in apparent contradiction with experiments. Indeed, experiments show that the shape of the droplet changes when the temperature is raised. More precisely, the facet radius slowly decreases for finally stabilizing after many hours (or even days) at a value that is a bit larger than that corresponding to the critical radius at the final temperature (given by Eq. (11a) and plotted in Fig. 6).

To raise the contradiction, let us study how the droplet shape evolves when the temperature is increased, with the additional constraint that its total height  $h_f$  cannot change. According to equation (5), this last condition reads

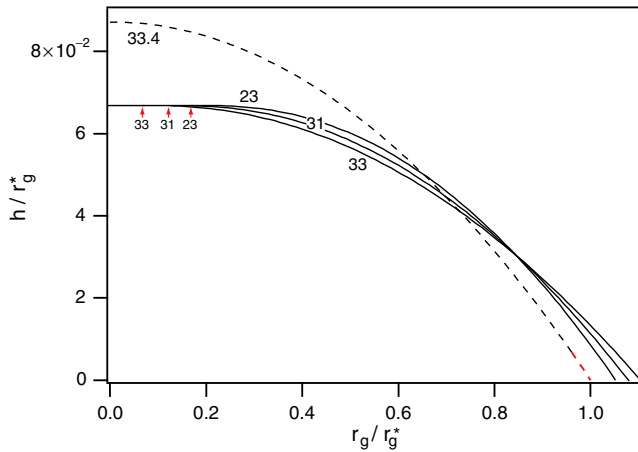
$$h_f = gr_g(k - 1 - \ln k) - \frac{p}{2} r_g^2 \left( \frac{k^2 - 1}{2} - k^2 \ln k \right), \quad (24)$$

where  $p$  is given by equation (9) as a function of the total volume  $V$  of the droplet (or, equivalently, as a function of  $r_g^*$ , the droplet radius in the nematic phase, see Eq. (20)). Replacing  $p$  by its expression in equation (24) leads to a third-degree equation in  $r_g$  which can be solved analytically using Mathematica. This procedure allows us to express  $r_g$  as a function of  $V$  (or  $r_g^*$ ),  $h_f$ , and  $k$ . The next





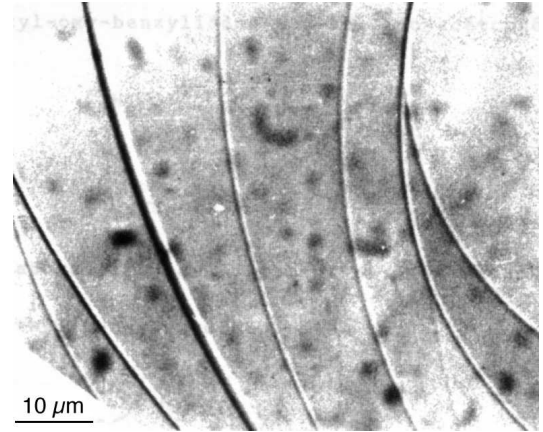
**Fig. 8.** Radius of the facet as a function of the temperature at increasing temperature (solid line, curve  $r_c^\dagger$ ). For comparison, we reported the critical radius as a function of the temperature (dashed line, curve  $r_c$ ).



**Fig. 9.** Evolution of the droplet profile when the temperature is increased. The temperature is indicated beside each profile and an arrow marks the edge of the facet. Although the droplet height does not change, the facet radius decreases. Because the volume is conserved, the droplet radius increases. The dashed line corresponds to the droplet profile in the nematic phase (spherical cap).

step is to replace  $r_g$  and  $p$  by their respective expressions in equation (7a). We thus obtain a new expression for the droplet energy  $e_o(k, s, g, r_g^*, h_f)$  at fixed height. The final step consists of finding the extrema of this function as a function of  $k$ . For each temperature  $T$ , this function has two extrema, the first one (at small  $k$ ) corresponding to the critical radius (with the droplet being marginally stable) and the other (at large  $k$ ) corresponding to a new “equilibrium” shape (it being agreed that  $h_f$  has been fixed).

The question now is to determine which of these two solutions the system chooses. At the beginning, the droplet is marginally stable (and, so, fairly far from its real equilibrium shape). So we propose as an ansatz that the droplet remains on this solution branch, *i.e.* it evolves while remaining marginally stable. Using this criterion, we can calculate the new shape of the droplet at the fi-



**Fig. 10.** Grandjean terraces separated by giant dislocations in a big droplet. Only a small part of the droplet is visible in this photograph taken in natural light (from Ref. [27]).

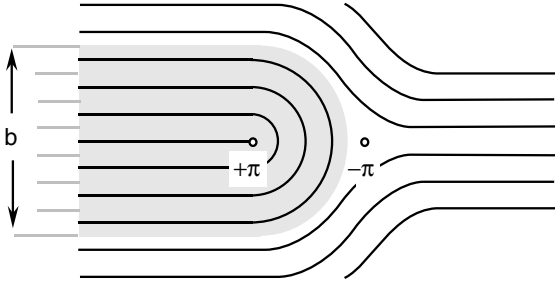
nal temperature. Figure 8 shows the new facet radius  $r_c^\dagger$  calculated this way. As expected, and in agreement with experiments,  $r_c^\dagger > r_c$ . Another prediction (which has not been checked experimentally) is that the droplet radius  $r_g$  must slightly increase, because the droplet approaches a spherical cap. This is visible on the profiles shown in Figure 9. Finally, this model predicts that the radius of curvature measured at the top of the droplet jumps at the transition temperature toward the nematic phase. This point would merit to be checked experimentally.

In conclusion, this approach of the problem could explain why the facet radius slowly decreases when the temperature is raised, in spite of the fact that no new dislocations nucleate. In the following section, we treat the problem of the stability of a droplet with respect to the formation of giant dislocations.

## 7 Stability with respect to the formation of giant dislocations

It was known for a long time that big droplets form Grandjean terraces [28]. The latter are separated by giant dislocations of very large Burgers vectors (many tens or even hundreds of layers) in the thick parts of the droplets. Such dislocations are shown in Figure 10.

This phenomenon is due to the tendency of the elementary dislocations to group together to reduce their core energy. Indeed, their elastic energy is proportional to their Burgers vector  $b$  (and not to  $b^2$  as in usual solids) [11, 29], a property that was checked experimentally only recently in films [30]. Thus, grouping  $n$  dislocations must be energetically favourable (at least in an infinite medium) because we do not change the total elastic energy, but we reduce the core energy (one core, or rather two, instead of  $n$ , if one considers the model shown in Fig. 11). This property was used to explain the grouping of dislocations in the menisci bordering free films [31] (in that case the two surfaces are free).



**Fig. 11.** Giant dislocation. The core is decomposed into a pair of  $(\pm\pi)$ -wedge disclination lines. These two lines have a singular core. This model was first proposed by Williams and Kléman [32].

In this section, we analyze by using the same method as in reference [31] in which conditions the dislocations group together in a smectic droplet. For the reasons explained before, we shall assume that the facet radius is equal to the critical radius and that the droplet profile is still given by equation (5) with  $c_f = 0$ .

Let us now consider a set of  $n$  elementary dislocations taken in the radius interval  $\rho \in [\delta - d/2, \delta + d/2]$  (with  $r_f < \delta < r_g$  and  $d \ll r_g$ ). The posed question is to determine in which condition it becomes preferable to group these  $n$  dislocations to form a giant dislocation of Burgers vector  $nb$  at the mean distance  $\delta$  from the droplet centre. To answer this question, let us determine the energy difference between the two possible configurations.

In the case of  $n$  elementary dislocations, the energy  $E_{ed}$  of the circular droplet slice of width  $d$  reads:

$$E_{ed} = 2\pi\delta \left( nE_d + \gamma \frac{[h'(\delta)]^2}{2} d \right). \quad (25)$$

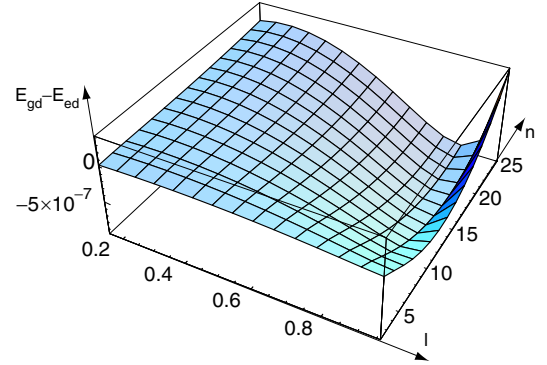
The first term represents the energy of the  $n$  elementary dislocations and the second the increase of surface energy due to the free surface inclination (with  $h'(\delta) = (dh(\rho)/d\rho)_{\rho=\delta}$ ). Because  $nb \approx -h'(\delta)d$ , the parameter  $d$  can be eliminated from equation (25) which becomes:

$$E_{ed} = 2\pi\delta n \left( E_d - \frac{\gamma}{2} bh'(\delta) \right). \quad (26)$$

For a giant dislocation of Burgers vector  $nb$  with the core structure described in Figure 11, the energy can be written in the form

$$E_{gd} = 2\pi\delta \left\{ 2E_c + \frac{nb\sqrt{KB}}{2} + \frac{\pi K}{2} \ln n + \frac{B\lambda n^2 b^2}{8\sqrt{2\pi\lambda h(\delta)}} [2A + (1 + A^{2/3})^{3/2}] \right\}. \quad (27)$$

In this expression,  $2E_c$  represents the core energy of the  $+\pi$  and  $-\pi$  disclinations (in the following, we shall take  $2E_c \approx E_d$ ). The second term corresponds to the elastic self-energy of the giant dislocation. The third term gives the curvature energy of the layers associated with the  $+\pi$



**Fig. 12.** Plot of the difference energy  $E_{gd} - E_{ed}$  as a function of  $n$  and  $l = \delta/r_g$  calculated for 8CB at  $T = 24^\circ\text{C}$ . In this graph,  $l_{min} \approx 0.39$  and  $n_{min} \approx 14$ , while  $l_{opt} \approx 0.87$  and  $n_{opt} \approx 16$ .

disclination. Finally, the last term corresponds to the elastic interaction energy between the giant dislocation and, both, the solid substrate and the free surface (for more details, see Ref. [22]). We recall that  $K$  and  $B$  are, respectively, the curvature and the compressibility moduli of the layers,  $\lambda = \sqrt{K/B}$  is the so-called penetration length, and  $A = (\gamma - \sqrt{KB})/(\gamma + \sqrt{KB})$  is a constant.

The next step is to compare the energy of the giant dislocation with the energy of the  $n$  elementary dislocations. We thus need to calculate  $E_{gd} - E_{ed}$  as a function of the free parameters  $n$  and  $\delta$ . This difference reads, according to equations (25) and (27),

$$E_{gd} - E_{ed} = 2\pi\delta \left\{ (1-n)E_d - n\frac{\gamma}{2}bh'(\delta) + \frac{nb\sqrt{KB}}{2\pi} + \frac{\pi K}{2} \ln n + \frac{B\lambda n^2 b^2}{8\sqrt{2\pi\lambda h(\delta)}} [2A + (1 + A^{2/3})^{3/2}] \right\}, \quad (28)$$

where  $h(\delta)$  and its derivative  $h'(\delta)$  can be calculated from equations (5) and (11a) (in the critical regime). Introducing the ratio  $l = \delta/r_g$  (with  $k < l < 1$ ), we calculate from these two equations

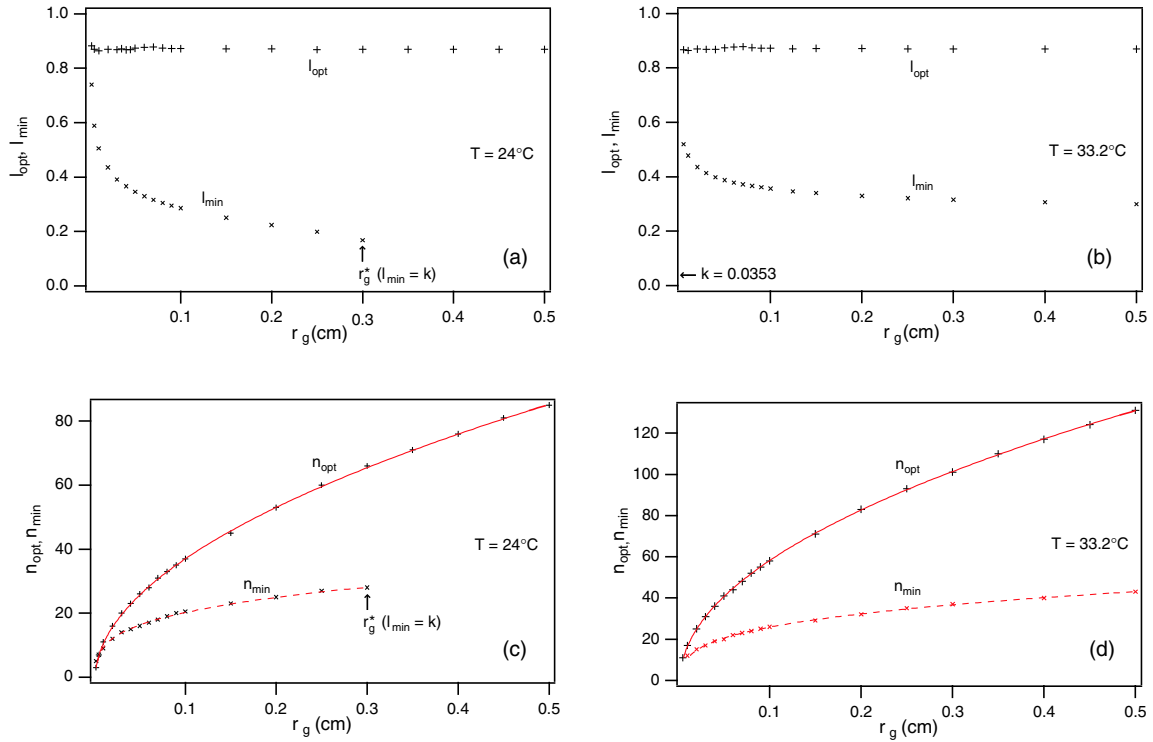
$$h(\delta)|_{\delta=lr_g} = \frac{g}{4k} [1 + 4k(l-1) - l^2 - 2k^2 \ln l] r_g \quad (28a)$$

and

$$h'(\delta)|_{\delta=lr_g} = -\frac{g(k-l)^2}{2kl}. \quad (28b)$$

Finally, we can express the energy difference  $E_{gd} - E_{ed}$  as a function of  $n$  and  $l$  for each temperature  $T$  and droplet radius  $r_g$ , knowing that  $E_d(\text{dyn}) = 2.52 \times 10^{-7} (33.4 - T)^{0.446}$  [26],  $B(\text{erg/cm}^3) \approx 0.665 \times 10^8 (33.4 - T)^{0.328}$ ,  $\lambda(\text{cm}) \approx 8.92 \times 10^{-8} (33.4 - T)^{-0.13}$  and  $K = B\lambda^2 \approx 7 \times 10^{-7} \text{ dyn}$  [11, 31]. The function  $E_{gd} - E_{ed}$  is plotted in Figure 12 by taking  $T = 24^\circ\text{C}$  and  $r_g = 300 \mu\text{m}$ .

This graph shows that  $E_{gd} - E_{ed}$  is positive at small values of  $l$ , but can become negative above some minimum value of  $l$  denoted by  $l_{min}$ . At the distance  $l_{min}r_g$  from the centre of the droplet, a giant dislocation of well-defined Burgers vector  $n_{min}$  (here, we measure the Burgers vector in number of layers, with  $n_{min} > 2$ ) may form



**Fig. 13.** a), b) Dimensionless distances  $l_{opt}$  and  $l_{min}$  as a function of the droplet size  $r_g$  calculated at two different temperatures; c), d) Burgers vectors (in number of layers)  $n_{opt}$  and  $n_{min}$  of the giant dislocations as a function of the droplet size  $r_g$  calculated for the same temperatures as in a) and b). Solid lines (respectively, dashed lines) are best fits to the numerical data of the type  $r_g^{0.5}$  (respectively,  $r_g^{0.3}$ ).

because it becomes energetically favourable. The quantity  $E_{gd} - E_{ed}$  has also an absolute minimum for some optimal values  $n_{opt}$  and  $l_{opt}$  of  $n$  and  $l$ . These quantities are plotted comparatively as a function of the droplet size  $r_g$  in Figure 13 at  $T = 24^\circ\text{C}$  and  $T = 33.2^\circ\text{C}$  (*i.e.*,  $0.2^\circ\text{C}$  below  $T_{NA} = 33.4^\circ\text{C}$ ), respectively.

We immediately note that  $l_{opt}$  is constant in the large range of radii explored ( $20\ \mu\text{m} < r_g < 5\ \text{mm}$ ) and is quasi-independent of the temperature chosen ( $l_{opt} \approx 0.87$  at  $T = 24^\circ\text{C}$  and  $l_{opt} \approx 0.89$  at  $T = 33.2^\circ\text{C}$ , Figs. 13a and b). This result shows that giant dislocations preferentially form close to the outer edge of the droplets. As for  $n_{opt}$ , it increases both with the temperature (at given radius) and with the droplet radius (at given temperature) as  $r_g^{1/2}$  (Figs. 13c and d). By contrast,  $l_{min}$  decreases when  $r_g$  increases, but the decrease is faster at low temperature than at high temperature (Figs. 13a and b). In addition, we note that  $l_{min}$  reaches its minimal possible value  $k$  for a droplet radius  $r_g^* \approx 3\ \text{mm}$  at  $T = 24^\circ\text{C}$  (Fig. 13a). This particular radius rapidly increases when the temperature increases and becomes so large close to the transition temperature that it does make sense to speak about it. In the same time,  $n_{min}$  increases with temperature (at a given radius) and with droplet radius as  $r_g^{0.3}$  (at a given temperature) (Figs. 13a and b).

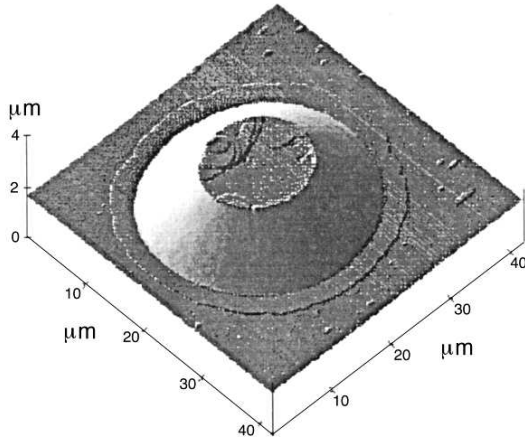
To summarize, giant dislocations form preferentially in the outer part of the droplet. Their Burgers vectors increase when the temperature increases in spite of the fact

that the curved part of the droplet surface adjacent to the facet is more stable at high temperature than at low temperature. For that reason, the whole droplet destabilizes only at low temperature if it is big enough (*i.e.*, if  $r_g > r_g^*$ ).

## 8 Concluding remarks

Our calculations lead to the conclusion that it is almost impossible to observe a droplet at equilibrium. Nevertheless, the shapes observed after cooling down small droplets from the nematic or the isotropic phase must be reproducible and self-similar, with a facet radius very close to the critical radius. This result reposes on the fact that the dislocation loops of smaller radius than  $r_c$  are unstable and, thus, rapidly collapse. This is clearly the case in small droplets in which the elastic force acting on dislocations (proportional to the overpressure  $\Delta P$ ) is large, being inversely proportional to  $r_g$ . On the contrary, shapes observed after a temperature increase change very slowly in time. In that case, the droplet height is blocked (the nucleation of new dislocations is forbidden), which does not prevent the facet from slowly decreasing in size. Our feeling is that the droplet evolves while remaining marginally stable. In that case, the final facet size must be larger than  $r_c$ , in agreement with experiments [19].

The problem is more complicated with big droplets because of the local formation of giant dislocations. The latter can form spontaneously at different places, resulting



**Fig. 14.** Three-dimensional view of a droplet of the chiral liquid crystal XI/12 obtained by tapping mode atomic force microscopy. The droplet has been deposited in the smectic A phase at high temperature (about 100 °C) and then slowly cooled down to the room temperature in the SmC\* phase (from Ref. [33]).

in various metastable shapes. This is perhaps one of the reasons why it was so difficult to observe faceted shapes with big droplets [19,20]. Another problem could come from wetting hysteresis. Indeed, we have implicitly assumed that the droplet radius can change easily in order to minimize the total energy, which excludes any hysteretic effects of the contact angle on the substrate. This is far from being the case in experiments, where hysteresis always exists. This can be a serious problem especially with big droplets in which the driving forces (inversely proportional to  $r_g^2$ ) are weak.

Finally, the model predicts that the curved part of the droplet matches tangentially the facet, on condition that the equilibrium equation (4) (which generalizes the Laplace equation) is globally satisfied. In that case, the profile is given by equation (5) with  $c_f = 0$  whatever the facet radius (always larger than the critical radius). The tangential matching is due to the fact that the dislocations repulse each other to reduce the excess of free surface energy. Indeed, the term in  $\gamma$  in equation (2) may be seen as a repulsive interaction term between the dislocations.

For this reason, the model cannot explain the angular matching observed in certain droplets of the liquid crystal 4O8 close to the smectic B phase transition [20]. In that case, which remains an exception, it is not excluded that the elastic constants  $K$  and  $B$  increase in such a way that the surface tension  $\gamma$  becomes smaller than  $\sqrt{KB}$ . The dislocations should then glide to the surface to form steps [22, 23] and the TLK model would become applicable. The explanation given in reference [21] based on finite-size effects and attractive elastic interactions between steps of the same sign via the solid substrate is thus perhaps the solution to this intriguing problem.

Let us still mention interesting unpublished observations by H. Dumoulin and P. Pieranski of smectic C\* droplets of a very viscous liquid crystal deposited on mica.

An example is shown in Figure 14. This image was obtained by using Tapping Mode Atomic Force Microscopy. In this droplet, the matching between the facet and the curved part of the surface is clearly angular (with an angle of 4.5° [33]). This result is very surprising because the elastic modulus  $B$  is usually smaller in SmC (or SmC\*) than in SmA due to the tilt of the molecules inside the layers [34]. For that reason, dislocations must be repelled from the free surface and our model should apply. One possibility for explaining the angle would be that the droplet has been “photographed” during the collapse of the upper dislocations, and so is strongly out of equilibrium. This is very possible inasmuch as steps are still visible on one side of the facet. In addition, the droplet is surrounded by a terrace of molecular height, which indicates that the liquid crystal wets mica and that the droplet is spreading (at least at high temperature). Thus, new and more controlled experiments are clearly necessary before to conclude about the existence or the non-existence of an angular matching at equilibrium in this material.

Finally, we emphasize that our model should apply as well to lamellar phases of diblock copolymers (like PS-PBMA). Indeed, the dislocations are strongly repulsed from the free surfaces in these systems, as was shown experimentally by electron microscopy [35]. Nevertheless, the ratio  $r_f/r_g$  must be smaller in copolymers than in usual smectic A because of the smallness of the ratio  $E_d/b \approx \sqrt{KB}$  (of the order of 1 dyn/cm [36]). As a consequence, facets should be more difficult to observe in copolymers than in usual smectic A.

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## References

1. G. Wulff, *Z. Kristallogr.* **34**, 449 (1901).
2. P. Nozières, *Shape and growth of crystals*, in *Solid Far From Equilibrium, Beg Rohu Lectures*, edited by C. Godreche (Cambridge University Press, Cambridge, 1992).
3. J.C. Heyraud, J.J. Métois, J.M. Bermond, *J. Cryst. Growth* **98**, 355 (1989).
4. J.M. Bermond, J.J. Métois, J.C. Heyraud, *Surf. Sci.* **416**, 430 (1998).
5. M.S. Hoogeman, M.A.J. Klik, D.C. Schlosser, L. Kuipers, J.W.M. Krenken, *Surf. Sci.* **448**, 142 (2000).
6. S. Balibar, H. Alles, A.Ya. Parshin, *Rev. Mod. Phys.* **77**, 317 (2005).
7. P. Pieranski, R. Barbet-Massin, P.E. Cladis, *Phys. Rev. A* **31**, 3912 (1985).
8. H. Stegemeyer, Th. Blümel, K. Hiltrop, H. Onusseit, F. Porsch, *Liq. Cryst.* **1**, 3 (1986).
9. P. Oswald, P. Pieranski, *Nematic and Cholesteric Liquid Crystals: Concepts and Physical Properties Illustrated by Experiments, The Liquid Crystal Book Series* (Taylor & Francis, 2005).
10. P. Oswald, F. Melo, C. Germain, *J. Phys. (Paris)* **50**, 3527 (1989).

11. P. Oswald, P. Pieranski, *Smectic and Columnar Liquid Crystals: Concepts and Physical Properties Illustrated by Experiments, The Liquid Crystal Book Series* (Taylor & Francis, 2005).
12. P. Pieranski, P. Sotta, D. Rohe, M. Imperor-Clerc, Phys. Rev. Lett. **84**, 2409 (2000).
13. P. Pieranski, L. Sittler, P. Sotta, M. Imperor-Clerc, Eur. J. Phys. E **5**, 317 (2001).
14. P. Pieranski, M. Bouchih, N. Ginestet, S. Popa-Nita, Eur. J. Phys. E **12**, 239 (2003).
15. P. Nozières, F. Pistolesi, S. Balibar, Eur. Phys. J. B **24**, 387 (2001).
16. J.D. Bernal, D. Crowfoot, Trans. Faraday Soc. **29**, 1032 (1933).
17. S. Chandrasekhar, Mol. Cryst. **2**, 71 (1966).
18. S. Chandrasekhar, N.V. Madhusudana, Acta Crystallogr., Sect. A **26**, 153 (1970).
19. J. Bechhoefer, P. Oswald, Europhys. Lett. **15**, 521 (1991).
20. J. Bechhoefer, L. Lejček, P. Oswald, J. Phys. II **2**, 27 (1992).
21. L. Lejček, J. Bechhoefer, P. Oswald, J. Phys. II **2**, 1511 (1992).
22. L. Lejček, P. Oswald, J. Phys. II **1**, 931 (1991).
23. This is due to the fact that the step energy per unit length, of the order of  $\gamma b$  (with  $\gamma$  the surface tension), is typically 3 to 10 times larger than the dislocation energy, of the order of  $\sqrt{KB}b$  (where  $K$  and  $B$  are, respectively, the curvature constant and the compressibility modulus of the layers). As a consequence, the dislocations are both repulsed by the solid and the free surfaces and place roughly at mid-distance between them (more precisely, at distance  $h/(1 + A^{3/2})$  from the solid surface [22], by denoting by  $h$  the local droplet thickness and by setting  $A = (\gamma - \sqrt{KB})/(\gamma + \sqrt{KB})$ ).
24. J.-C. Gémard, R. Holyst, P. Oswald, Phys. Rev. Lett. **78**, 1924 (1997).
25. R. Jaquet, F. Schneider, Phys. Rev. E **67**, 021707 (2003).
26. A. Zywockinski, F. Picano, P. Oswald, J.C. Gémard, Phys. Rev. E **62**, 8133 (2000).
27. C. Williams, *Défauts de structure dans les smectiques A*, D.Sc. Thesis, University of Paris XI, Orsay (1976).
28. F. Grandjean, Bull. Soc. Fr. Minéral. **39**, 164 (1916).
29. M. Kléman, J. Phys. (Paris) **35**, 595 (1974).
30. J.C. Gémard, C. Laroche, P. Oswald, Phys. Rev. E **58**, 5923 (1998).
31. F. Picano, R. Holyst, P. Oswald, Phys. Rev. E **62**, 3747 (2000).
32. C.E. Williams, M. Kléman, J. Phys. (Paris) Colloq. **36**, C1-315 (1976).
33. H. Dumoulin, *Etude d'interfaces de cristaux liquides par microscopie de proximité*, PhD Thesis, University of Paris XI, Orsay (1996).
34. S. Shibahara, J. Yamamoto, Y. Takanishi, K. Ishikawa, H. Takezoe, J. Phys. Soc. Jpn. **71**, 802 (2002).
35. M. Maaloum, D. Ausserré, D. Chatenay, G. Coulon, Y. Gallot, Phys. Rev. Lett. **68**, 1575 (1992).
36. M.S. Turner, M. Maaloum, D. Ausserré, J.-F. Joanny, M. Kunz, J. Phys. II **4**, 689 (1994).